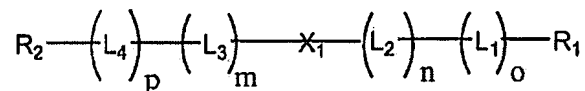


Listing of Claims

1. (currently amended) An oligonucleotide prodrug of the formula (I):



(I)

wherein:

R_1 and R_2 are independently H or a polymer residue;

L_1 and L_4 are independently selected releasable linking moieties;

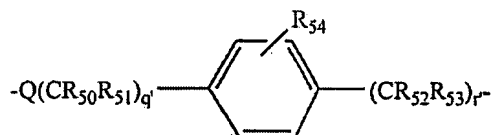
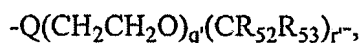
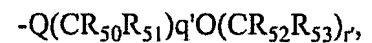
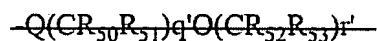
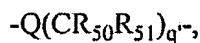
L_2 and L_3 are independently selected bifunctional spacing groups each comprising from about 2 to about 10 carbon atoms;

X_1 is a single or double stranded nucleotide or an oligonucleotide residue

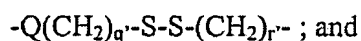
wherein the oligonucleotide ranges in size from 10 to 1,000 nucleotides;

m , n , o and p are independently zero or a positive integer, provided that either $(o + n)$ or $(p + m) \geq 2$; and

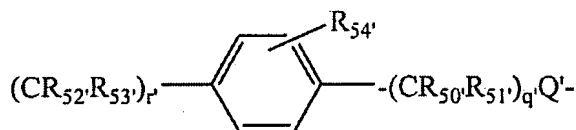
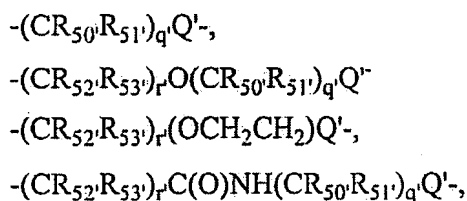
wherein L_3 is selected from the group consisting of:



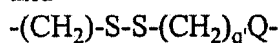
and



L_2 is selected from the group consisting of:



and



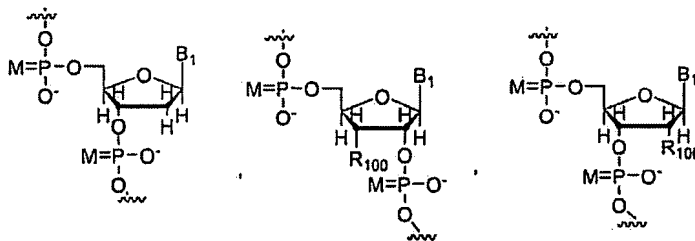
wherein.

Q and Q' are independently selected from O, S or NH;

R₅₀₋₅₃ and R_{50'-53'} are independently selected from the group consisting of hydrogen, C₁₋₆ alkyls, C₃₋₁₂ branched alkyls, C₃₋₈ cycloalkyls, C₁₋₆ substituted alkyls, C₃₋₈ substituted cyloalkyls, aryls substituted aryls, aralkyls, C₁₋₆ heteroalkyls, substituted C₁₋₆ heteroalkyls, C₁₋₆ alkoxy, phenoxy and C₁₋₆ heteroalkoxy;

R₅₄ and R_{54'} are independently selected from the group consisting of hydrogen, C₁₋₆ alkyls, C₃₋₁₂ branched alkyls, C₃₋₈ cycloalkyls, C₁₋₆ substituted alkyls, C₃₋₈ substituted cycloalkyls, aryls substituted aryls, aralkyls, C₁₋₆ heteroalkyls, substituted C₁₋₆ heteroalkyls, C₁₋₆ alkoxy, phenoxy, C₁₋₆ heteroalkoxy, NO₂, haloalkyl and halogen; and q' and r' are each a positive integer.

2. (currently amended) The prodrug of claim 1, wherein ~~said nucleotide~~ the oligonucleotide comprises a nucleotide that is selected from the group consisting of



wherein

M is O or S;

B₁ and B₂ are independently selected from the group consisting of A (adenine), G (guanine), C (cytosine), T (thymine), U (uracil) and modified bases; R₁₀₀ and R₁₀₁ are independently selected from the group consisting of H, OR' where R' is H, a C₁₋₆ alkyl, substituted alkyls, nitro, halo and aryl

3. (currently amended) The prodrug of claim 1, wherein said oligonucleotide ~~[[is]]~~ contains from about 10 to about 1000 nucleotides.

4. (currently amended) The prodrug of claim ~~[[1]]~~ 2, wherein M is S.

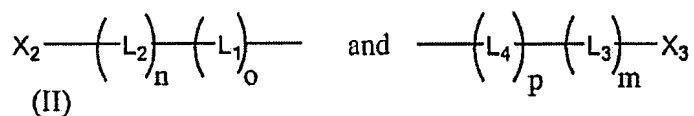
5. (currently amended) The prodrug of claim 1, wherein the oligonucleotide residue is a ~~phosphorothioate~~ phosphorothioate oligonucleotide residue.

6. (previously presented) The prodrug of claim 1, wherein said oligonucleotide residue is an antisense oligonucleotide residue or oligodeoxynucleotide residue.

7. (previously presented) The prodrug of claim 6, wherein said antisense oligonucleotide residue or oligodeoxynucleotide residue is selected from the group consisting of, oligonucleotides and oligodeoxynucleotides with phosphodiester backbones or phosphorothioate backbones, LNA, PNA, tricyclo-DNA, decoy ODN, ribozymes, spiegelmers, and CpG oligomers.

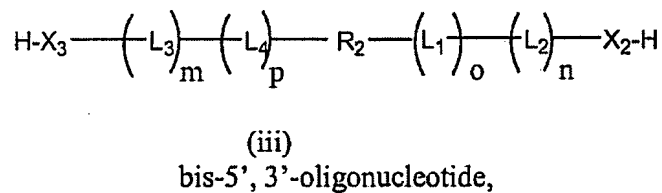
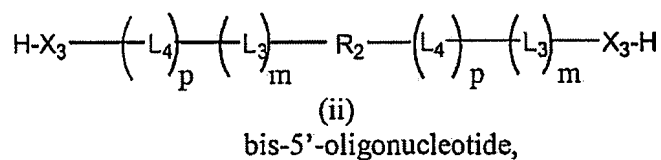
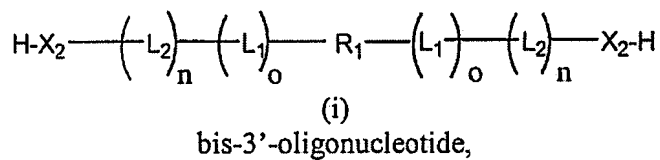
8. (withdrawn) The prodrug of claim 6, wherein said antisense oligonucleotide has a sequence selected from the group consisting of SEQ ID NO: 1, SEQ ID NO: 2, SEQ ID NO: 3, and SEQ ID NO: 4, wherein n of SEQ ID NO: 4 is any compatible nucleotide.

9. (previously presented) The prodrug of claim 1, wherein at least one of R₁ and R₂ is a polymeric residue having a capping group A, selected from the group consisting of OH, NH₂, SH, CO₂H, C₁₋₆ alkyls,

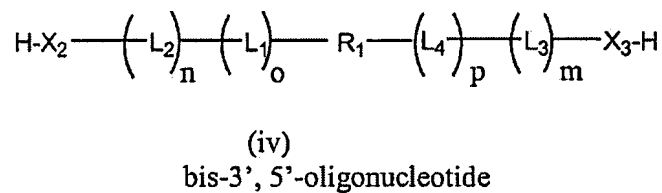


wherein X_2 and X_3 are independently selected nucleotide or oligonucleotide residues.

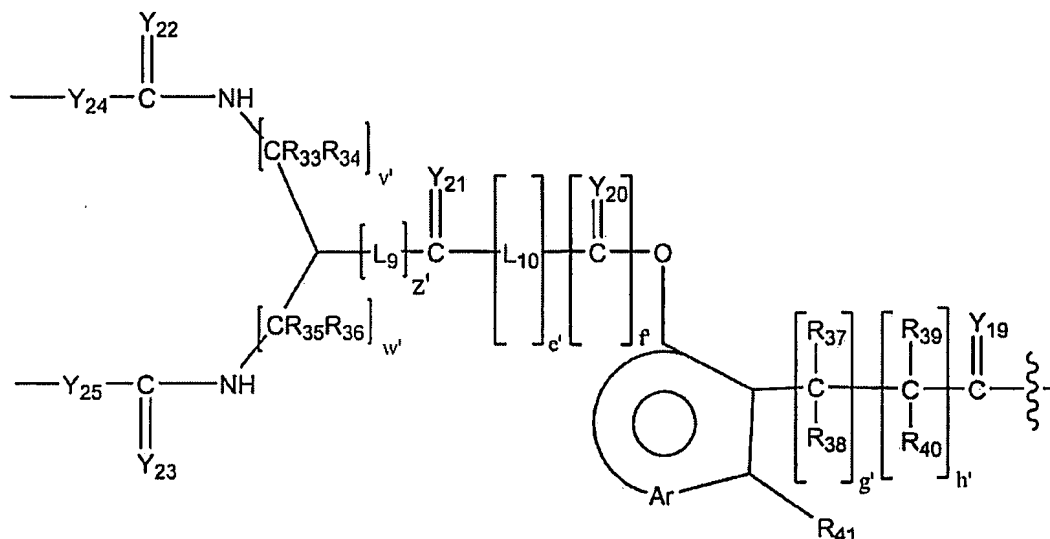
10. (previously presented) A prodrug of claim 9, selected from the group consisting of:



and



11. (currently amended) The prodrug of claim 1 wherein L_4 is selected from the group consisting of:



wherein:

Y_{1-25} are independently selected from the group consisting of O, S or NR_9 ;

R_{6-7} , R_{9-13} , R_{16-25} , and R_{27-41} are independently selected from the group consisting of hydrogen, C_{1-6} alkyls, C_{3-12} branched alkyls, C_{3-8} cycloalkyls, C_{1-6} substituted alkyls, C_{3-8} substituted cycloalkyls, aryls, substituted aryls, aralkyls, C_{1-6} heteroalkyls, substituted C_{1-6} heteroalkyls, C_{1-6} alkoxy, phenoxy and C_{1-6} heteroalkoxy;

Ar is a moiety which forms a multi-substituted aromatic hydrocarbon or a multi-substituted heterocyclic group;

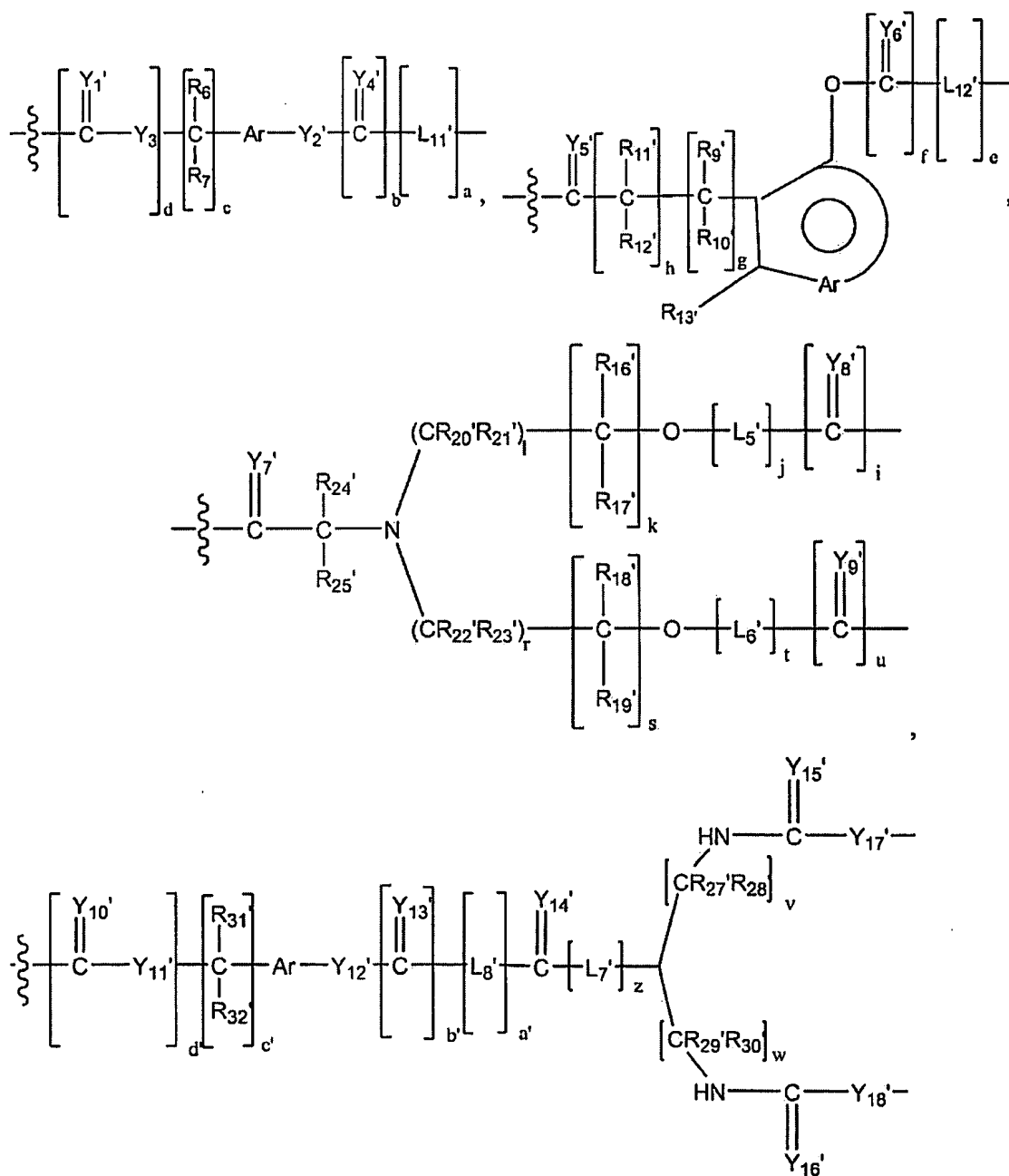
L_{5-12} are independently selected bifunctional spacers;

c, h, k, l, r, s, v, w, v', w', c', and h' are independently selected positive integers;

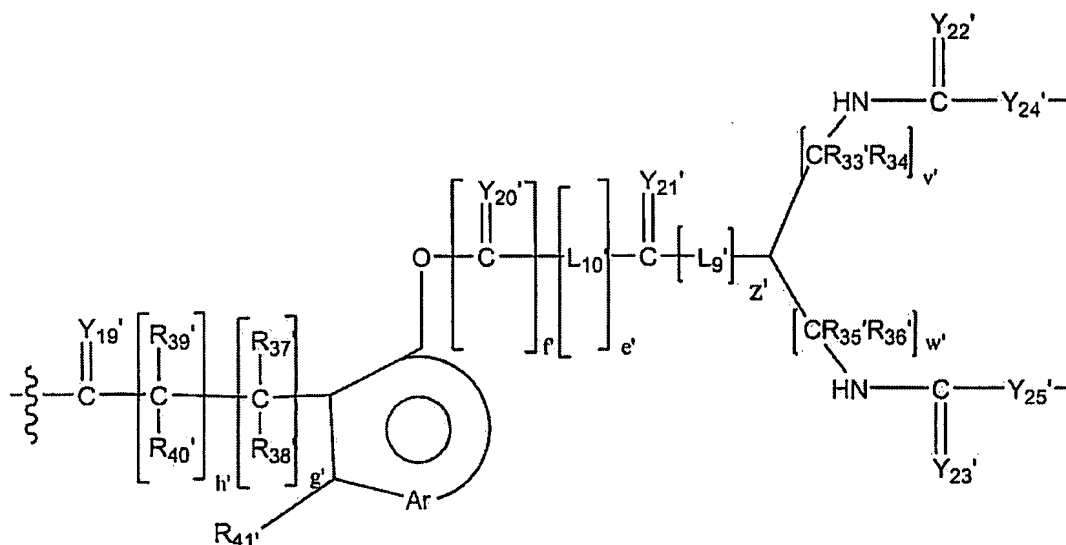
a, e, g, j, t, z, a', z', e' and g' are independently either zero or a positive integer; and

b, d, f, i, u, [[q,]] b', d' and f' are independently zero or one.

12. (currently amended) The prodrug of claim 1 wherein L_1 is selected from the group consisting of:



and



wherein,

Y_1, Y_{25} , are independently selected from the group consisting of O, S or NR_9 ;

R_{6-7} , R_{9-13} , R_{16-25} , and R_{27-41} are independently selected from the group consisting of hydrogen, C_{1-6} alkyls, C_{3-12} branched alkyls, C_{3-8} cycloalkyls, C_{1-6} substituted alkyls, C_{3-8} substituted cycloalkyls, aryls, substituted aryls, aralkyls, C_{1-6} heteroalkyls, substituted C_{1-6} heteroalkyls, C_{1-6} alkoxy, phenoxy and C_{1-6} heteroalkoxy; and

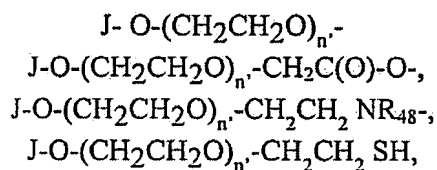
L_{5-12} are independently selected bifunctional spacers;

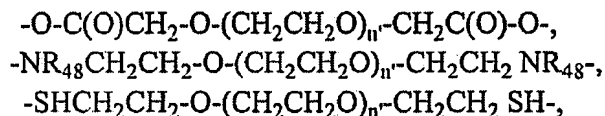
c, h, k, l, r, s, v, w, v', w', c', and h' are independently selected positive integers;

$a, e, g, i, t, z, a', z', e'$ and g' are independently either zero or a positive integer; and

b, d, f, i, u, b', d' and f' are independently zero or one.

13. (previously presented) The prodrug of claim 1 wherein R_{1-2} are each polyalkylene oxides.
14. (previously presented) The prodrug of claim 1 wherein R_{1-2} are each polyethylene glycols.
15. (previously presented) The prodrug of claim 1 wherein R_{1-2} are independently selected from the group consisting of:





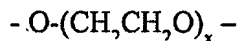
wherein

n' is the degree of polymerization;

R_{48} is selected from the group consisting of hydrogen, C_{1-6} alkyls, C_{3-12} branched alkyls, C_{3-8} cycloalkyls, C_{1-6} substituted alkyls, C_{3-8} substituted cycloalkyls, aryls substituted aryls, aralkyls, C_{1-6} heteroalkyls, substituted C_{1-6} heteroalkyls, C_{1-6} alkoxy, phenoxy and C_{1-6} heteroalkoxy; and

J is a capping group.

16. (previously presented) The prodrug of claim 1, wherein R_{1-2} are independently



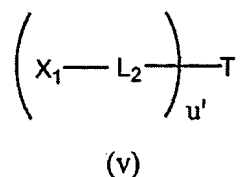
wherein x is a positive integer selected so that the weight average molecular weight is at least about 2,000 Da to about 136,000 Da.

17. (previously presented) The prodrug of claim 1, wherein R_{1-2} independently have a weight average molecular weight of from about 3,000 Da to about 100,000 Da.

18. (previously presented) The prodrug of claim 1, wherein R_{1-2} independently have a weight average molecular weight of from about 5,000 Da to about 40,000 Da.

19. (previously presented) The prodrug of claim 8, wherein said antisense oligonucleotide is oblimersen (SEQ ID NO: 1).

20. (withdrawn) An oligonucleotide prodrug of the formula:



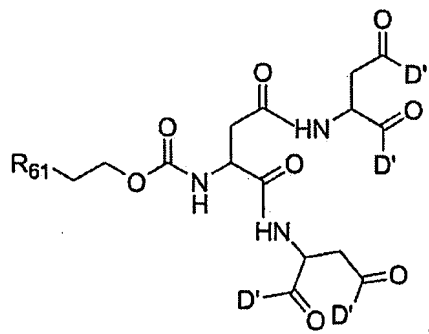
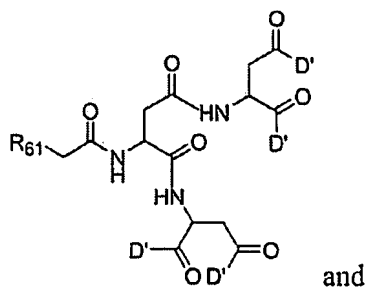
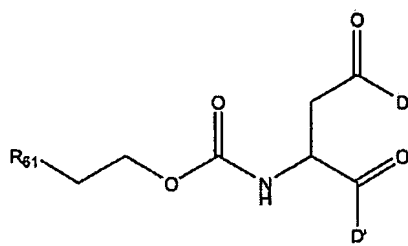
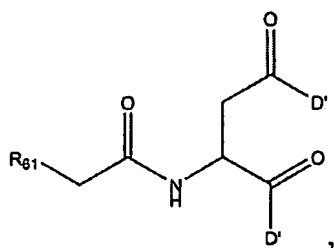
wherein:

L_2 is a spacing group;

X_1 is a nucleotide or an oligonucleotide residue;

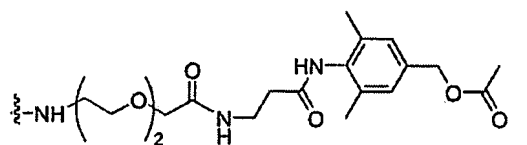
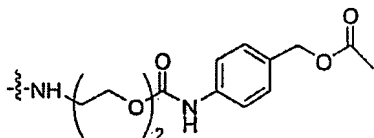
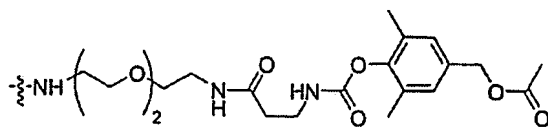
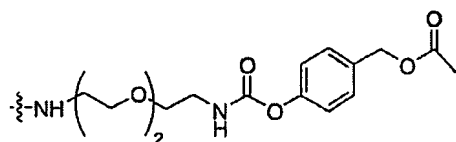
u' is a positive integer; and

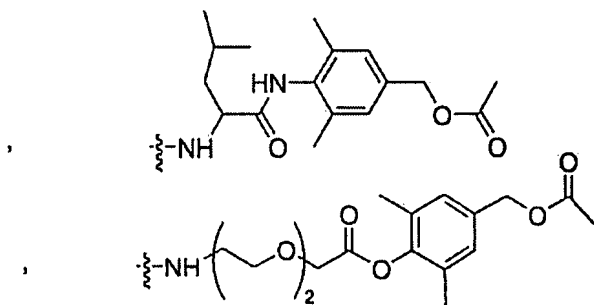
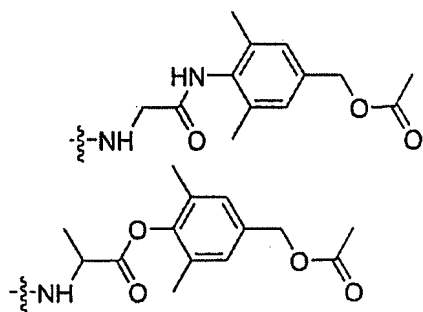
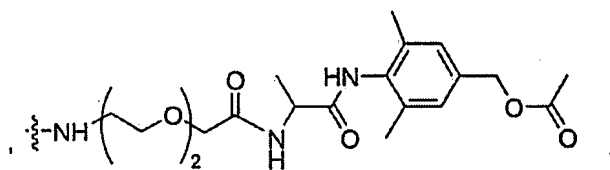
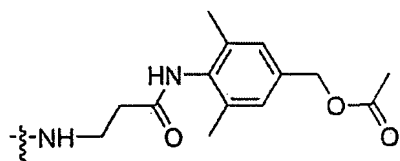
T is a member of the group consisting of:

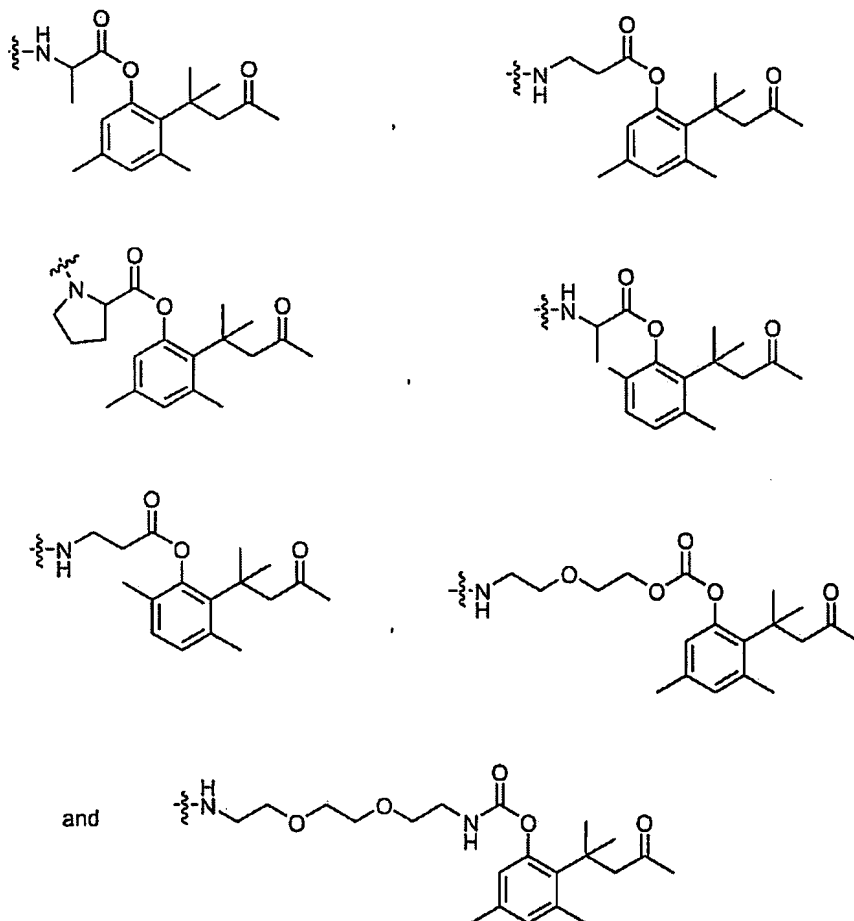


wherein:

D' is one of

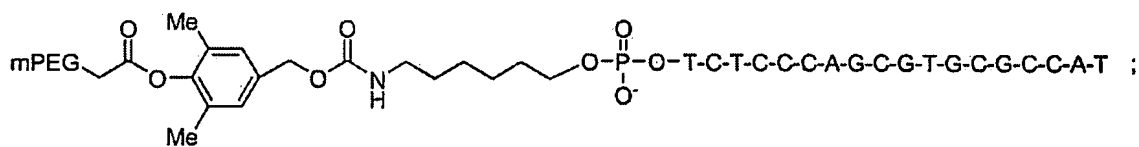
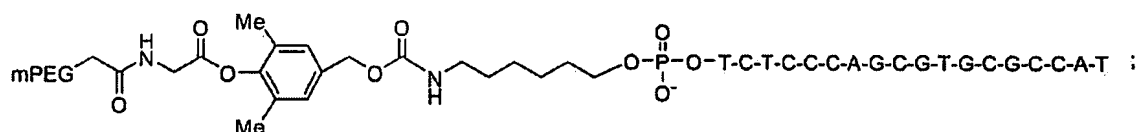




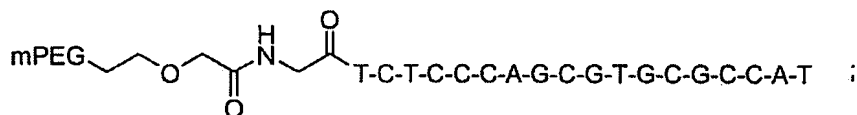


and wherein R_{61} is a polymer residue.

21. (previously presented) A compound of claim 1 selected from the group consisting of:



and



all of which comprise an oligonucleotide of SEQ ID NO: 1.

22. (withdrawn) A method of making a prodrug comprising:

reacting a compound of the formula:

R_2 - L_4 -leaving group

with a compound of the formula:

H - L_3 - X_1

under conditions sufficient to form a prodrug of the formula

R_2 - L_4 - L_3 - X_1 ,

wherein:

R_2 is a polymer residue;

L_4 is a releasable linking moiety;

L_3 is a spacing group;

X_1 is a nucleotide or an oligonucleotide residue.

23. (withdrawn) A method of treating a mammal, comprising administering to a mammal in need of such treatment an effective amount of a compound of claim 1.

24. (withdrawn) The method of claim 23, wherein the mammal is being treated for cancer.

25. (withdrawn) The method of claim 23, wherein X_1 is an antisense oligonucleotide.

26. (withdrawn) The method of claim 23 wherein the mammal is also treated with a second anticancer agent that is administered simultaneously or sequentially with the oligonucleotide prodrug.